

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	0	(514/275andhpparsactivators). CCLS	US-PGPUB	OR	OFF	2006/02/02 15:14

Patent Search

STN Full Text

Number(s): JP 2000-369891

0 answer(s) in PCTFULL

Select a database to search for equivalents for EP / US / WO full text.

☐ CPlus



☒ INPADOC



☐ WPIDS/WPINDEX



Click "Search" to search the selected database.

Search

* * * * * Welcome to STN International * * * * *

NEWS 1 Web Page URLs for STN Seminar Schedule - N. America
NEWS 2 "Ask CAS" for self-help around the clock
NEWS 3 DEC 05 CASREACT(R) - Over 10 million reactions available
NEWS 4 DEC 14 2006 MeSH terms loaded in MEDLINE/LMEDLINE
NEWS 5 DEC 14 2006 MeSH terms loaded for MEDLINE file segment of TOXCENTER
NEWS 6 DEC 14 CA/CAPLUS to be enhanced with updated IPC codes
NEWS 7 DEC 21 IPC search and display fields enhanced in CA/CAPLUS with the
IPC reform
NEWS 8 DEC 23 New IPC8 SEARCH, DISPLAY, and SELECT fields in USPATFULL/
USPAT2
NEWS 9 JAN 13 IPC 8 searching in IFIPAT, IFIUDB, and IFICDB
NEWS 10 JAN 13 New IPC 8 SEARCH, DISPLAY, and SELECT enhancements added to
INPADOC
NEWS 11 JAN 17 Pre-1988 INPI data added to MARPAT
NEWS 12 JAN 17 IPC 8 in the WPI family of databases including WPIFV
NEWS 13 JAN 30 Saved answer limit increased
NEWS 14 JAN 31 Monthly current-awareness alert (SDI) frequency
added to TULSA

NEWS EXPRESS JANUARY 03 CURRENT VERSION FOR WINDOWS IS V8.01,
CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
AND CURRENT DISCOVER FILE IS DATED 19 DECEMBER 2005.
V8.0 USERS CAN OBTAIN THE UPGRADE TO V8.01 AT
<http://download.cas.org/express/v8.0-Discover/>

NEWS HOURS STN Operating Hours Plus Help Desk Availability
NEWS INTER General Internet Information
NEWS LOGIN Welcome Banner and News Items
NEWS PHONE Direct Dial and Telecommunication Network Access to STN
NEWS WWW CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that specific topic.

All use of STN is subject to the provisions of the STN Customer agreement. Please note that this agreement limits use to scientific research. Use for software development or design or implementation of commercial gateways or other similar uses is prohibited and may result in loss of user privileges and other penalties.

* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 11:26:48 ON 02 FEB 2006

=> file reg

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

FILE 'REGISTRY' ENTERED AT 11:26:55 ON 02 FEB 2006

USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.

PLEASE SEE "HELP USAGETERMS" FOR DETAILS.

COPYRIGHT (C) 2006 American Chemical Society (ACS)

Property values tagged with IC are from the ZIC/VINITI data file provided by InfoChem.

STRUCTURE FILE UPDATES: 31 JAN 2006 HIGHEST RN 873191-05-0
 DICTIONARY FILE UPDATES: 31 JAN 2006 HIGHEST RN 873191-05-0

New CAS Information Use Policies, enter HELP USAGETERMS for details.

TSCA INFORMATION NOW CURRENT THROUGH JULY 14, 2005

Please note that search-term pricing does apply when
 conducting SmartSELECT searches.

```
*****
*
* The CA roles and document type information have been removed from *
* the IDE default display format and the ED field has been added,   *
* effective March 20, 2005. A new display format, IDERL, is now    *
* available and contains the CA role and document type information. *
*
*****
```

Structure search iteration limits have been increased. See HELP SLIMITS
 for details.

REGISTRY includes numerically searchable data for experimental and
 predicted properties as well as tags indicating availability of
 experimental property data in the original document. For information
 on property searching in REGISTRY, refer to:

<http://www.cas.org/ONLINE/UG/regprops.html>

=>

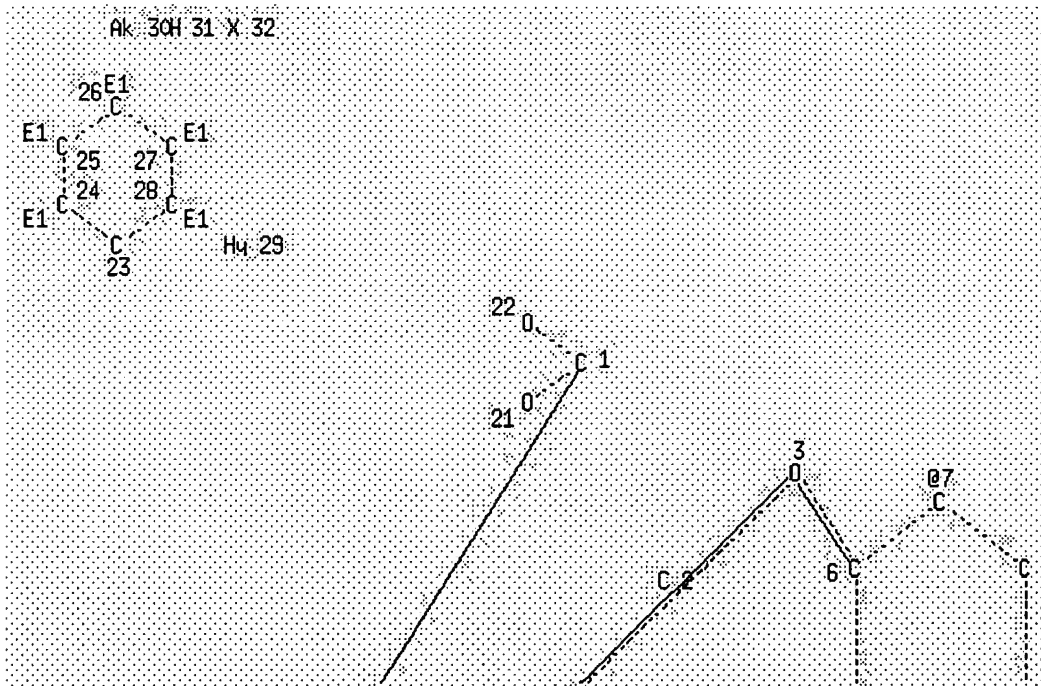
Uploading structure

L1 STRUCTURE UPLOADED

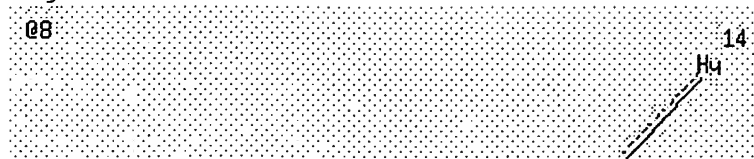
=> d 11

L1 HAS NO ANSWERS

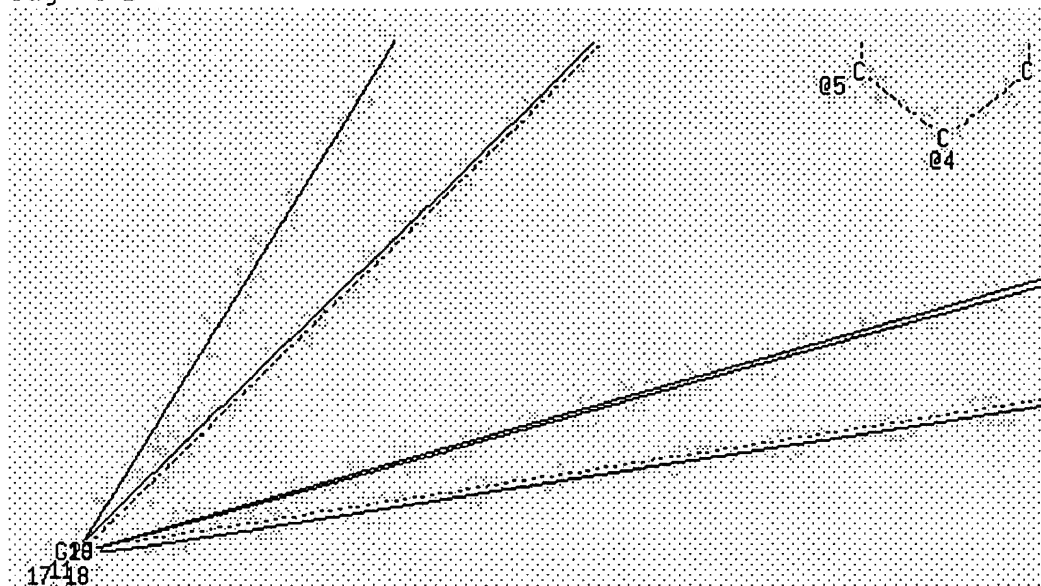
L1 STR



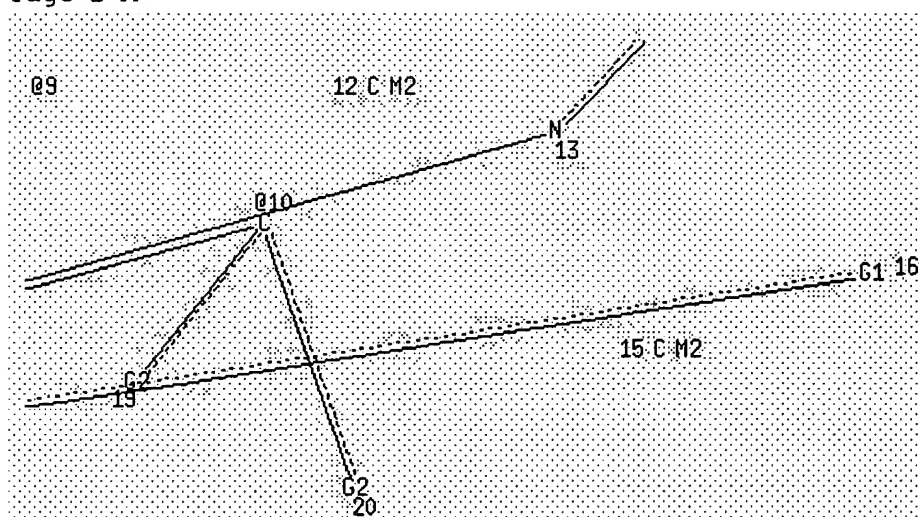
Page 1-A



Page 1-B



Page 2-A



Page 2-B

```

VAR G1=23/29
VAR G2=30/31/32
REP G18=(0-3) 12-10 12-13
REP G19=(1-3) 15-13 15-16
REP G20=(1-2) 2-1 2-3
VPA 10-4/5/7/8/9 S
NODE ATTRIBUTES:
HCOUNT IS M2 AT 12
HCOUNT IS M2 AT 15
HCOUNT IS E1 AT 24
HCOUNT IS E1 AT 25
HCOUNT IS E1 AT 26
HCOUNT IS E1 AT 27
HCOUNT IS E1 AT 28

```

```

NSPEC  IS C      AT   1
NSPEC  IS C      AT   2
NSPEC  IS C      AT   3
NSPEC  IS R      AT   4
NSPEC  IS R      AT   5
NSPEC  IS R      AT   6
NSPEC  IS R      AT   7
NSPEC  IS R      AT   8
NSPEC  IS R      AT   9
NSPEC  IS C      AT  10
NSPEC  IS C      AT  11
NSPEC  IS C      AT  12
NSPEC  IS C      AT  13
NSPEC  IS C      AT  14
NSPEC  IS C      AT  15
NSPEC  IS C      AT  16
NSPEC  IS C      AT  17
NSPEC  IS C      AT  18
NSPEC  IS C      AT  19
NSPEC  IS C      AT  20
NSPEC  IS C      AT  21
NSPEC  IS C      AT  22

```

DEFAULT MLEVEL IS ATOM

```

MLEVEL  IS CLASS AT   1   2   3  10  12  13  15  21  22  23  24  25  26  27  28  29  30
      31  32

```

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RSPEC I

NUMBER OF NODES IS 32

STEREO ATTRIBUTES: NONE

=> s 11

SAMPLE SEARCH INITIATED 11:30:54 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 3326 TO ITERATE

60.1% PROCESSED 2000 ITERATIONS

0 ANSWERS

INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 63062 TO 69978

PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s 11 full

THE ESTIMATED SEARCH COST FOR FILE 'REGISTRY' IS 166.50 U.S. DOLLARS

DO YOU WANT TO CONTINUE WITH THIS REQUEST? (Y)/N or END:y

FULL SEARCH INITIATED 11:30:58 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 67878 TO ITERATE

100.0% PROCESSED 67878 ITERATIONS

26 ANSWERS

SEARCH TIME: 00.00.01

L3 26 SEA SSS FUL L1

```
=> file hcaplus
COST IN U.S. DOLLARS          SINCE FILE      TOTAL
                                ENTRY      SESSION
FULL ESTIMATED COST          169.58      169.79
```

FILE 'HCAPLUS' ENTERED AT 11:31:02 ON 02 FEB 2006
 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
 PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
 COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)

Copyright of the articles to which records in this database refer is held by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications. The CA Lexicon is the copyrighted intellectual property of the the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

FILE COVERS 1907 - 2 Feb 2006 VOL 144 ISS 6
 FILE LAST UPDATED: 1 Feb 2006 (20060201/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

```
=> s l3
L4          5 L3

=> s l4 and seiichiro, m?/au
          1 SEIICHIRO, M?/AU
L5          0 L4 AND SEIICHIRO, M?/AU

=> s l4 and yoshida, s?/au
          8841 YOSHIDA, S?/AU
L6          1 L4 AND YOSHIDA, S?/AU

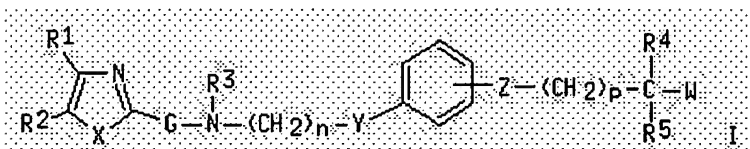
=> d l6, ibib abs hitstr, 1
```

L6 ANSWER 1 OF 1 HCAPLUS COPYRIGHT 2006 ACS on STN

Full Text	Citing References
ACCESSION NUMBER:	2002:449667 HCAPLUS
DOCUMENT NUMBER:	137:20370
TITLE:	Preparation of thiazole and benzothiazole derivatives as activators of PPAR (peroxisome proliferator activated receptor)
INVENTOR(S):	Yoshida, Shinichi; Sakuma, Shogo; Endo, Tsuyoshi; Tendo, Atsushi; Takahashi, Toshihiro; Kobayashi, Kunio; Mochiduki, Nobutaka; Yamakawa, Tomio; Kanda, Takashi; Masui, Seiichiro
PATENT ASSIGNEE(S):	Nippon Chemiphar Co., Ltd., Japan
SOURCE:	PCT Int. Appl., 79 pp. CODEN: PIXXD2
DOCUMENT TYPE:	Patent
LANGUAGE:	Japanese
FAMILY ACC. NUM. COUNT:	1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002046176	A1	20020613	WO 2001-JP10577	20011204
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2002024139	A5	20020618	AU 2002-24139	20011204
PRIORITY APPLN. INFO.:			JP 2000-369891	A 20001205
			WO 2001-JP10577	W 20011204
OTHER SOURCE(S):		MARPAT 137:20370		
GI				



AB The title compds. I [R1, R2 = H, halo, etc.; R1 and R2 together with the carbon atoms to which they are bonded may form a benzene ring; X = S, etc.; G = carbonyl, etc.; R3 = alkyl, etc.; Y = CH2, etc.; Z = O, etc.; n = 0 - 5; p = 0 - 5; R4, R5 = H, alkyl; W = carboxyl, etc.] are prepd. In a test for activation of PPAR α , compds. of this invention in vitro showed EC50 values of 0.2 μ M to 2.5 μ M. In a test for activation of PPAR γ , compds. of this invention in vitro showed EC50 values of 0.7 μ M to 5 μ M. In a test for activation of PPAR δ , compds. of this invention in vitro showed EC50 values of 0.9 μ M to 4.4 μ M.

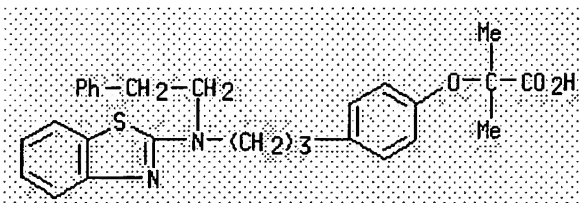
IT 435339-90-5P 435339-91-6P 435339-92-7P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of thiazole and benzothiazole derivs. as activators of peroxisome proliferator activated receptor)

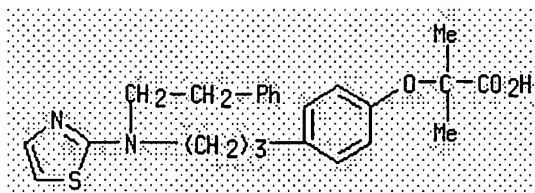
RN 435339-90-5 HCAPLUS

CN Propanoic acid, 2-[4-[3-[2-benzothiazolyl(2-phenylethyl)amino]propyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



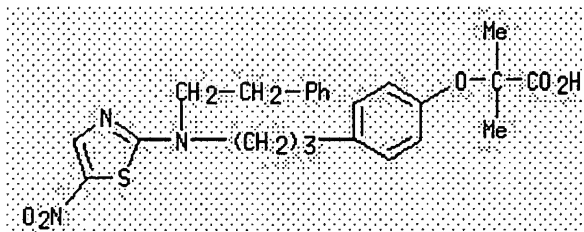
RN 435339-91-6 HCAPLUS

CN Propanoic acid, 2-methyl-2-[4-[3-[(2-phenylethyl)-2-thiazolylamino]propyl]phenoxy]- (9CI) (CA INDEX NAME)



RN 435339-92-7 HCAPLUS

CN Propanoic acid, 2-methyl-2-[4-[3-[(5-nitro-2-thiazolyl)(2-phenylethyl)amino]propyl]phenoxy]- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 49 THERE ARE 49 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> fil PCTFULL;s (JP 2000-369891)/pn,apps

COST IN U.S. DOLLARS

SINCE FILE ENTRY	TOTAL SESSION
------------------	---------------

FULL ESTIMATED COST

10.17	179.96
-------	--------

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE ENTRY	TOTAL SESSION
------------------	---------------

CA SUBSCRIBER PRICE

-0.75	-0.75
-------	-------

FILE 'PCTFULL' ENTERED AT 11:32:03 ON 02 FEB 2006

COPYRIGHT (C) 2006 Univentio

FILE LAST UPDATED: 3 JAN 2006 <20060103/UP>

MOST RECENT UPDATE WEEK: 200552 <200552/EW>

FILE COVERS 1978 TO DATE

>>> IMAGES ARE AVAILABLE ONLINE AND FOR EMAIL-PRINTS <<<

>>> UPDATING DELAYED DUE TO DELIVERY FORMAT CHANGES. <<<

>>> NEW IPC8 DATA AND FUNCTIONALITY NOT YET AVAILABLE IN THIS FILE.
USE IPC7 FORMAT FOR SEARCHING THE IPC. WATCH THIS SPACE FOR FURTHER
DEVELOPMENTS AND SEE OUR NEWS SECTION FOR FURTHER INFORMATION
ABOUT THE IPC REFORM <<<

0 (JP 2000-369891)/PN
(JP2000369891/PN)

0 (JP 2000-369891)/APPS
(JP2000-369891/APPS)

L7 0 (JP 2000-369891)/PN,APPS

=> d his

=> s 14 not 16

COMBINATION OF STRUCTURE AND TEXT TERMS NOT VALID

The query entered contains both search terms created by

structure-building or screen commands and text search terms. L#s created via the STRUCTURE or SCREEN commands must be searched in the structures files separately from text terms or profiles. The L# answer sets from structure searches can be used in crossover searches and can be combined with text terms.

=> file hcaplus

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	2.32	193.37
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	0.00	-0.75

FILE 'HCAPLUS' ENTERED AT 11:33:54 ON 02 FEB 2006
 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT.
 PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
 COPYRIGHT (C) 2006 AMERICAN CHEMICAL SOCIETY (ACS)

Copyright of the articles to which records in this database refer is held by the publishers listed in the PUBLISHER (PB) field (available for records published or updated in Chemical Abstracts after December 26, 1996), unless otherwise indicated in the original publications. The CA Lexicon is the copyrighted intellectual property of the the American Chemical Society and is provided to assist you in searching databases on STN. Any dissemination, distribution, copying, or storing of this information, without the prior written consent of CAS, is strictly prohibited.

FILE COVERS 1907 - 2 Feb 2006 VOL 144 ISS 6
 FILE LAST UPDATED: 1 Feb 2006 (20060201/ED)

New CAS Information Use Policies, enter HELP USAGETERMS for details.

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> d his

(FILE 'HOME' ENTERED AT 11:26:48 ON 02 FEB 2006)

FILE 'REGISTRY' ENTERED AT 11:26:55 ON 02 FEB 2006

L1 STRUCTURE UPLOADED
 L2 0 S L1
 L3 26 S L1 FULL

FILE 'HCAPLUS' ENTERED AT 11:31:02 ON 02 FEB 2006

L4 5 S L3
 L5 0 S L4 AND SEIICHIRO, M?/AU
 L6 1 S L4 AND YOSHIDA, S?/AU

FILE 'PCTFULL' ENTERED AT 11:32:03 ON 02 FEB 2006

L7 0 S (JP 2000-369891)/PN,APPS

FILE 'CAPLUS' ENTERED AT 11:32:22 ON 02 FEB 2006

L8 1 S (JP 2000-369891)/PN,APPS
 SEL PRN

FILE 'EPFULL' ENTERED AT 11:32:22 ON 02 FEB 2006
L9 1 S E1-2/APPS

FILE 'USPATFULL' ENTERED AT 11:32:24 ON 02 FEB 2006
L10 0 S E1-2/APPS

FILE 'PCTFULL' ENTERED AT 11:32:25 ON 02 FEB 2006
L11 1 S E1-2/APPS

FILE 'HCAPLUS' ENTERED AT 11:33:54 ON 02 FEB 2006

=> s 14 not 16

L12 4 L4 NOT L6

=> d 112, ibib abs hitstr, 1-4

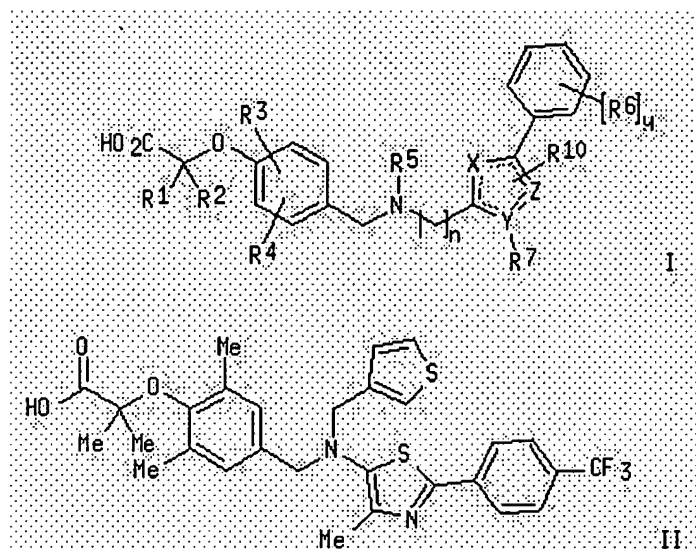
L12 ANSWER 1 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN

Full
Text

ACCESSION NUMBER: 2004:2833 HCAPLUS
DOCUMENT NUMBER: 140:77141
TITLE: Preparation of 2-[4-(heteroarylaminomethyl)phenoxy]-2-methylpropanoates for treating a hPPAR mediated diseases
INVENTOR(S): Dodic, Nerina; Dumaitre, Bernard Andre; Gellibert, Francoise Jeanne; Sierra, Michael Lawrence
PATENT ASSIGNEE(S): Smithkline Beecham Corporation, USA
SOURCE: PCT Int. Appl., 89 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
<u>WO 2004000785</u>	A2	20031231	<u>WO 2003-EP6417</u>	20030618
<u>WO 2004000785</u>	A3	20041014		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
<u>EP 1513796</u>	A2	20050316	<u>EP 2003-735642</u>	20030618
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
<u>JP 2005529965</u>	T2	20051006	<u>JP 2004-514763</u>	20030618
<u>US 2005222424</u>	A1	20051006	<u>US 2004-518347</u>	20041217
<u>PRIORITY APPLN. INFO.:</u>			<u>GB 2002-14139</u>	A 20020619
			<u>WO 2003-EP6417</u>	W 20030618

OTHER SOURCE(S): MARPAT 140:77141
GI



AB The title compds. [I; R1, R2 = H, alkyl; R3, R4 = H, alkyl, OMe, CF₃, allyl, halo; n = 0-1; at least of X, Z and Y = O, S, N; R6 = alkyl, CF₃, OMe, OCF₃, halo; y = 0-5; R7 = H, CF₃, alkyl (optionally substituted by phenyl), alkenyl with the proviso that when Z = S, O, R7 = H; R10 = H, alkyl; R5 = H, alkyl, alkoxyalkyl, alkenyl, alkoxy, etc.], useful for treatment of a hPPAR disease or condition such as dyslipidemia, syndrome X, heart failure, hypercholesterolemia, cardiovascular disease, diabetes, insulin resistance, hyperlipidemia, obesity, anorexia bulimia and anorexia nervosa (no biol. data given), were prepd. Thus, reacting Et 2-(4-bromomethyl-2,6-dimethylphenoxy)-2-methylpropionate with [4-methyl-2-(4-trifluoromethylphenyl)thiazol-5-yl]thiophen-3-ylmethylamine (preps. given) in the presence of cesium carbonate in 3-methyl-2-butanone followed by hydrolysis afforded II. Pharmaceutical compn. comprising the compd. I.

IT 639783-41-8P 639783-43-0P 639783-51-0P

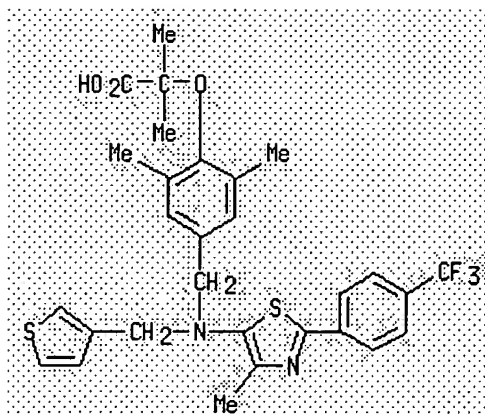
639784-00-2P 639784-02-4P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of 2-[4-(heteroarylaminomethylphenoxy)]-2-methylpropanoates for treating a hPPAR mediated diseases)

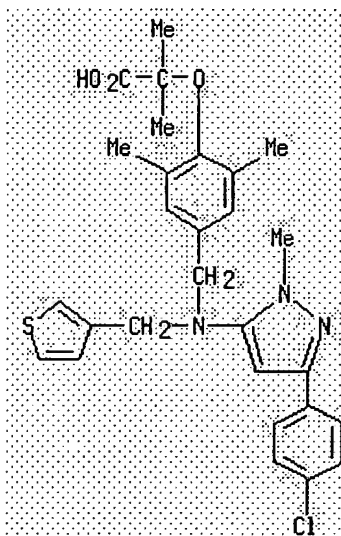
RN 639783-41-8 HCAPLUS

CN Propanoic acid, 2-[2,6-dimethyl-4-[[[4-methyl-2-[4-(trifluoromethyl)phenyl]-5-thiazolyl](3-thienylmethyl)amino]methyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



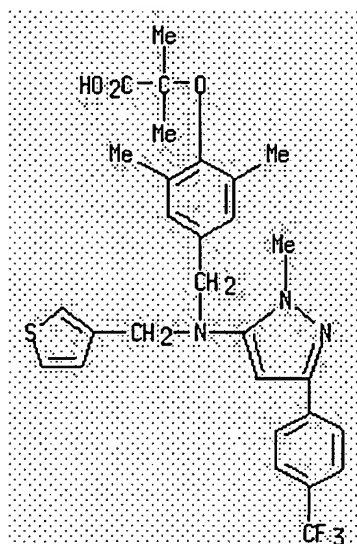
RN 639783-43-0 HCAPLUS

CN Propanoic acid, 2-[4-[[[3-(4-chlorophenyl)-1-methyl-1H-pyrazol-5-yl](3-thienylmethyl)amino]methyl]-2,6-dimethylphenoxy]-2-methyl- (9CI) (CA INDEX NAME)



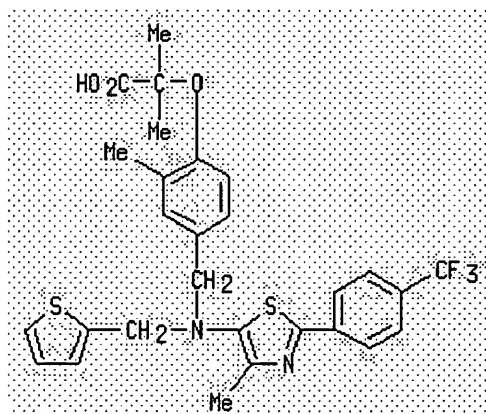
RN 639783-51-0 HCAPLUS

CN Propanoic acid, 2-[2,6-dimethyl-4-[[[1-methyl-3-[4-(trifluoromethyl)phenyl]-1H-pyrazol-5-yl](3-thienylmethyl)amino]methyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



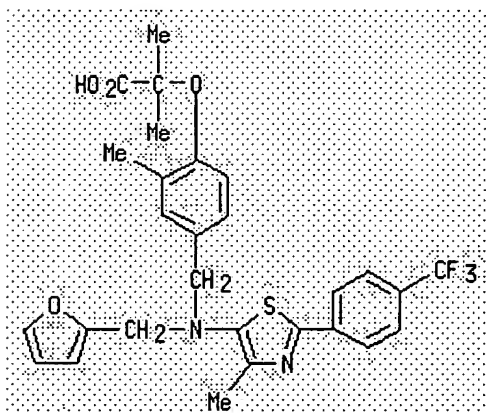
RN 639784-00-2 HCAPLUS

CN Propanoic acid, 2-methyl-2-[2-methyl-4-[[[4-methyl-2-[4-(trifluoromethyl)phenyl]-5-thiazolyl](2-thienylmethyl)amino]methyl]phenoxy]- (9CI) (CA INDEX NAME)



RN 639784-02-4 HCAPLUS

CN Propanoic acid, 2-[4-[[[2-furanylmethyl][4-methyl-2-[4-(trifluoromethyl)phenyl]-5-thiazolyl]amino]methyl]-2-methylphenoxy]-2-methyl- (9CI) (CA INDEX NAME)



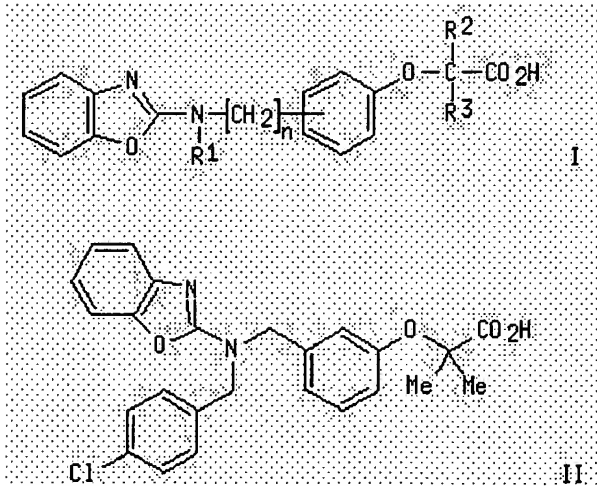
L12 ANSWER 2 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN



ACCESSION NUMBER: 2003:922669 HCAPLUS
 DOCUMENT NUMBER: 139:395923
 TITLE: Preparation of benzoxazoles as PPAR α agonists
 INVENTOR(S): Yamazaki, Yukiyo; Toma, Tsutomu; Nishikawa, Masahiro; Ozawa, Hidefumi; Okuda, Ayumu; Abe, Kazutoyo; Oda, Soichi
 PATENT ASSIGNEE(S): Kowa Co., Ltd., Japan
 SOURCE: U.S., 63 pp.
 CODEN: USXXAM
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 6653334	B1	20031125	US 2002-329547	20021227
JP 2004210776	A2	20040729	JP 2003-428197	20031224
EP 1433786	A1	20040630	EP 2003-29917	20031229

R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK
 PRIORITY APPLN. INFO.: US 2002-329547 A 20021227
 OTHER SOURCE(S): MARPAT 139:395923
 GI



AB The title compds. [I; R1 = H, alkyl, arylalkyl, etc.; R2, R3 = H, Me, Et; n = 1-3] and their salts, which selectively activate PPAR α , and are useful in preventing and/or treating hyperlipidemia, arteriosclerosis, diabetes, inflammation and heart diseases, were prepd. E.g., a 4-step synthesis of II (starting from 3-hydroxybenzaldehyde and Et 2-bromoisobutyrate) which showed EC₅₀ of 0.001 μ M, 0.2 μ M and >10 μ M with respect to hPPAR α , hPPAR γ and hPPAR δ , resp., was given. Pharmaceutical compn. comprising the compd. I is claimed.

IT 627095-30-1P 627095-45-8P 627095-57-2P
627095-63-0P 627095-70-9P 627095-78-7P
627095-99-2P 627096-10-0P 627096-58-6P
627096-63-3P 627096-73-5P 627096-87-1P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

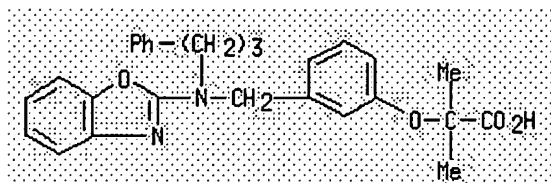
```



```

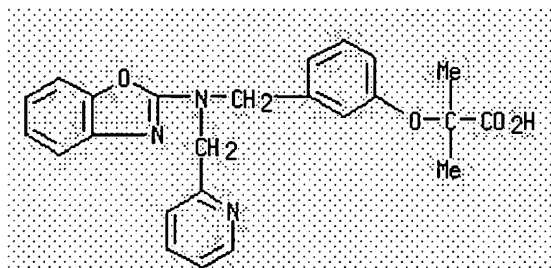
RN 627095-30-1 HCAPLUS

CN	Propanoic acid, 2-[3-[[2-benzoxazolyl (3-phenylpropyl) amino]methyl]phenoxy]-2-methyl- (9CI)	(CA INDEX NAME)
----	--	-----------------



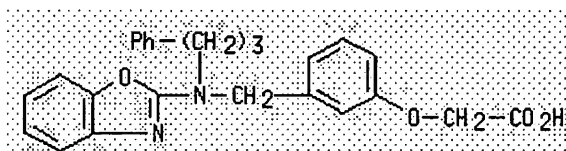
RN 627095-45-8 HCAPLUS

CN Propanoic acid, 2-[3-[[2-benzoxazolyl(2-pyridinylmethyl)amino]methyl]pheno
xy]-2-methyl- (9CI) (CA INDEX NAME)



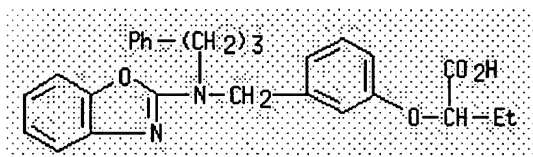
RN 627095-57-2 HCAPLUS

CN Acetic acid, [3-[[2-benzoxazolyl(3-phenylpropyl)amino]methyl]phenoxy]-
(9CI) (CA INDEX NAME)



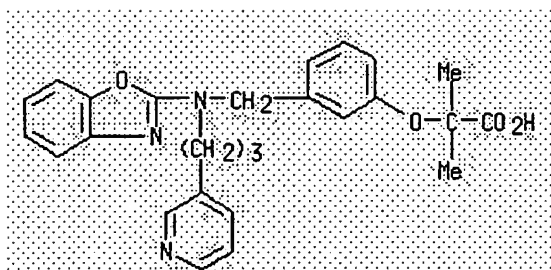
RN 627095-63-0 HCAPLUS

CN Butanoic acid, 2-[3-[[2-benzoxazolyl(3-phenylpropyl)amino]methyl]phenoxy]-
(9CI) (CA INDEX NAME)



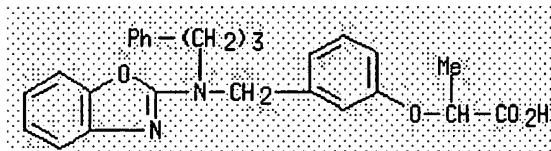
RN 627095-70-9 HCAPLUS

CN Propanoic acid, 2-[3-[[2-benzoxazolyl[3-(3-pyridinyl)propyl]amino]methyl]p
henoxy]-2-methyl- (9CI) (CA INDEX NAME)



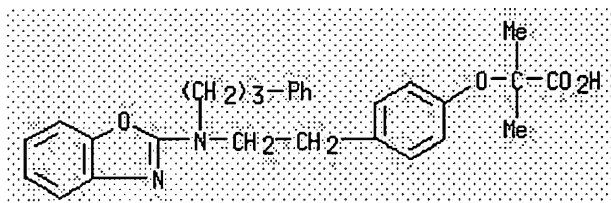
RN 627095-78-7 HCAPLUS

CN Propanoic acid, 2-[3-[[2-benzoxazolyl(3-phenylpropyl)amino]methyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



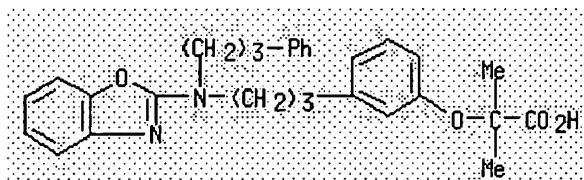
RN 627095-99-2 HCAPLUS

CN Propanoic acid, 2-[4-[[2-[2-benzoxazolyl(3-phenylpropyl)amino]ethyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



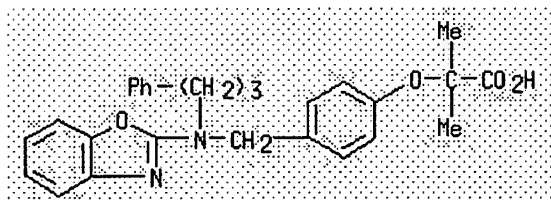
RN 627096-10-0 HCAPLUS

CN Propanoic acid, 2-[3-[3-[2-benzoxazolyl(3-phenylpropyl)amino]propyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



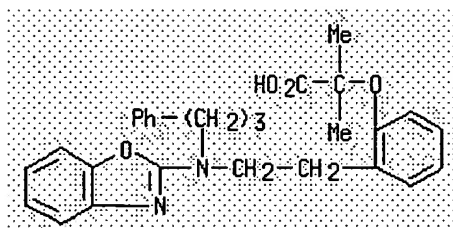
RN 627096-58-6 HCAPLUS

CN Propanoic acid, 2-[4-[[2-benzoxazolyl(3-phenylpropyl)amino]methyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



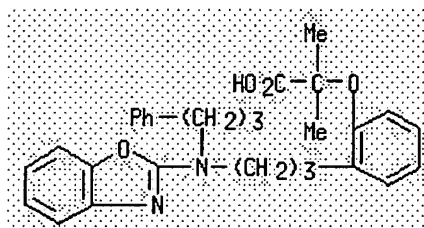
RN 627096-63-3 HCAPLUS

CN Propanoic acid, 2-[2-[2-[2-benzoxazolyl(3-phenylpropyl)amino]ethyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



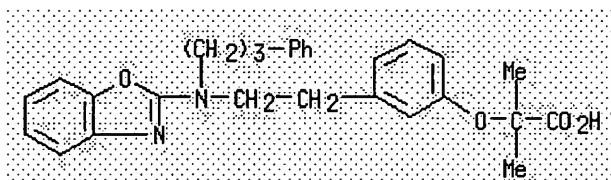
RN 627096-73-5 HCAPLUS

CN Propanoic acid, 2-[2-[3-[2-benzoxazolyl(3-phenylpropyl)amino]propyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



RN 627096-87-1 HCAPLUS

CN Propanoic acid, 2-[3-[2-[2-benzoxazolyl(3-phenylpropyl)amino]ethyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 30 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L12 ANSWER 3 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN

Full Text	References
-----------	------------

ACCESSION NUMBER: 2003:719457 HCAPLUS

DOCUMENT NUMBER: 139:245779

TITLE: Preparation of phenoxyalkanoic acid derivatives as hPPAR activators for treatment of diabetes and cardiovascular diseases

INVENTOR(S): Cadilla, Rodolfo; Henke, Brad Richard; Lambert, Millard H., III; Liu, Guangcheng Kevin; Smith, Jennifer Susan

PATENT ASSIGNEE(S): Smithkline Beecham Corporation, USA

SOURCE: PCT Int. Appl., 174 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

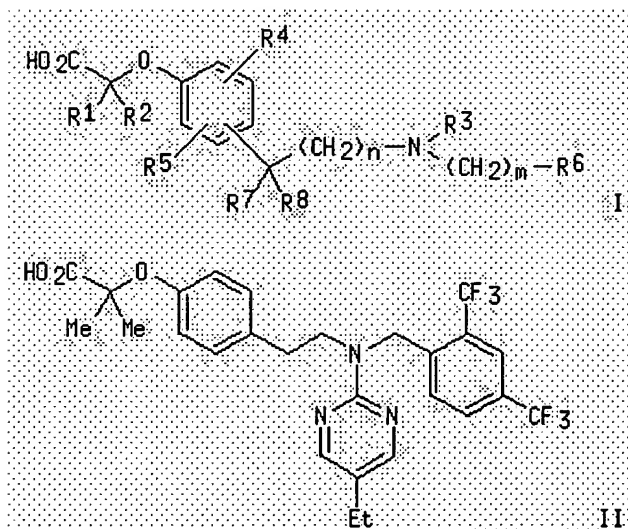
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003074495	A1	20030912	WO 2003-US5953	20030225
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,				

PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ,
 UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW
 RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY,
 KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES,
 FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, SE, SI, SK, TR, BF,
 BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

<u>AU 2003224632</u>	A1	20030916	<u>AU 2003-224632</u>	20030225
<u>EP 1480957</u>	A1	20041201	<u>EP 2003-721310</u>	20030225
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
<u>US 2005137212</u>	A1	20050623	<u>US 2003-505333</u>	20030225
<u>JP 2005532272</u>	T2	20051027	<u>JP 2003-572964</u>	20030225
<u>PRIORITY APPLN. INFO.:</u>			<u>US 2002-360975P</u>	P 20020301
			<u>WO 2003-US5953</u>	W 20030225

OTHER SOURCE(S): MARPAT 139:245779
 GI



AB Title compds. I [wherein R1 and R2 = independently H, F, CF₃, or alkyl; or CR1R2 = cycloalkyl; R3 = (un)substituted heteroaryl; R4 and R5 = independently H, (perfluoro)alkyl, (perfluoro)alkoxy, halo, or CN; R6 = (un)substituted Ph or heteroaryl; R7 and R8 = independently H, F, CF₃, or alkyl with the proviso that the C to which R7 and R8 are bonded is either meta or para to the depicted O; m and n = independently 1-2; or pharmaceutically acceptable salts, solvates, acid isosteres, or hydrolyzable esters thereof] were prepd. as human peroxisome proliferator activated receptor (hPPAR) activators (no data). For example, Me 2-[4-[2-[[2,4-bis(trifluoromethyl)benzyl]amino]ethyl]phenoxy]-2-methylpropanoate was coupled with 2-chloro-5-ethylpyrimidine using DIEA in toluene to give the tertiary amine (38%). Hydrolysis of the ester with NaOH provided II (48%). Methods for treating diseases or conditions assocd. with hPPAR α , hPPAR γ , or hPPAR δ , such as diabetes and cardiovascular diseases, comprising administration of a therapeutically effective amt. of I or a pharmaceutical compn. comprising I are also disclosed (no data).

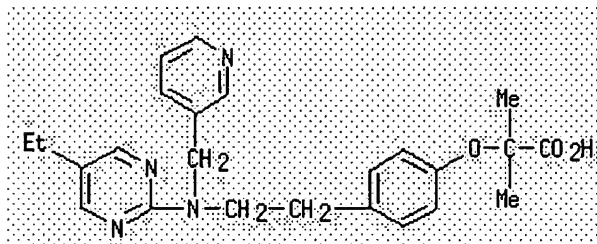
IT 596114-46-4P, 2-[4-[2-[(5-Ethylpyrimidin-2-yl)(pyridin-3-ylmethyl)amino]ethyl]phenoxy]-2-methylpropanoic acid 596114-49-7P, 2-[4-[2-[(5-Isopropylpyrimidin-2-yl)(pyridin-3-ylmethyl)amino]ethyl]phenoxy]-2-methylpropanoic acid 596114-52-2P

, 2-[4-[2-[(5-Ethylpyrimidin-2-yl) (pyridin-4-ylmethyl) amino]ethyl]phenoxy]-2-methylpropanoic acid **596114-86-2P**, 2-[4-[2-[Benzyl (5-ethylpyrimidin-2-yl) amino]ethyl]phenoxy]-2-methylpropanoic acid
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(PPAR activator; prepn. of phenoxyalkanoic acid derivs. as hPPAR activators for treatment of diabetes, cardiovascular diseases, and other disorders)

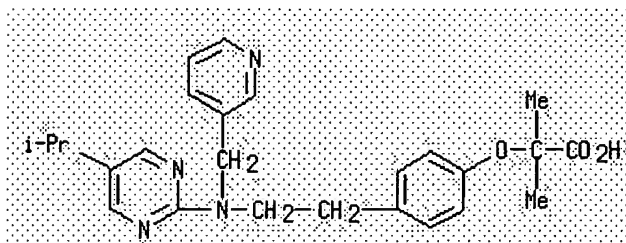
RN **596114-46-4** HCAPLUS

CN Propanoic acid, 2-[4-[2-[(5-ethyl-2-pyrimidinyl) (3-pyridinylmethyl) amino]ethyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



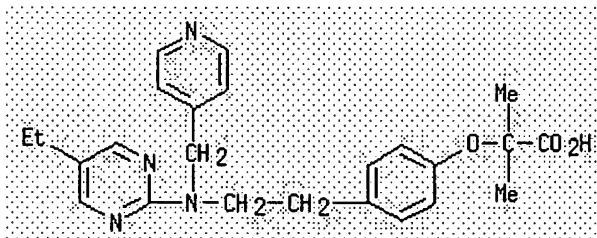
RN **596114-49-7** HCAPLUS

CN Propanoic acid, 2-methyl-2-[4-[2-[[5-(1-methylethyl)-2-pyrimidinyl] (3-pyridinylmethyl) amino]ethyl]phenoxy]- (9CI) (CA INDEX NAME)



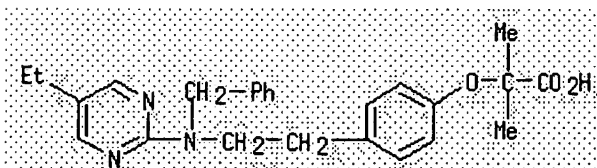
RN **596114-52-2** HCAPLUS

CN Propanoic acid, 2-[4-[2-[(5-ethyl-2-pyrimidinyl) (4-pyridinylmethyl) amino]ethyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



RN **596114-86-2** HCAPLUS

CN Propanoic acid, 2-[4-[2-[(5-ethyl-2-pyrimidinyl) (phenylmethyl) amino]ethyl]phenoxy]-2-methyl- (9CI) (CA INDEX NAME)



REFERENCE COUNT:

10

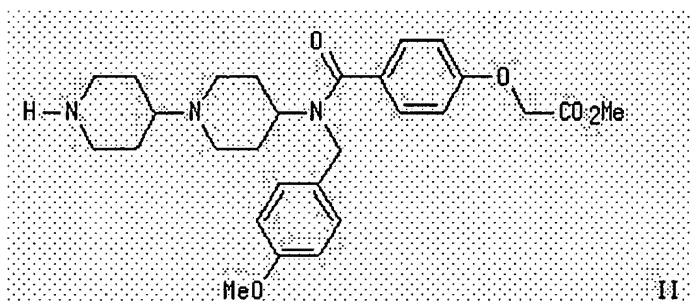
THERE ARE 10 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L12 ANSWER 4 OF 4 HCAPLUS COPYRIGHT 2006 ACS on STN

Full Text	Single References
--------------	----------------------

ACCESSION NUMBER: 1998:259658 HCAPLUS
 DOCUMENT NUMBER: 128:294701
 TITLE: Preparation of N-bipiperidinybenzamides and analogs
 as cell adhesion inhibitors
 INVENTOR(S): Pieper, Helmut; Linz, Guenter; Austel, Volkhard;
 Himmelsbach, Frank; Guth, Brian; Weisenberger,
 Johannes
 PATENT ASSIGNEE(S): Dr. Karl Thomae G.m.b.H., Germany
 SOURCE: Ger. Offen., 40 pp.
 CODEN: GWXXBX
 DOCUMENT TYPE: Patent
 LANGUAGE: German
 FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
<u>DE 19643331</u>	A1	19980423	<u>DE 1996-19643331</u>	19961021
<u>WO 9817646</u>	A1	19980430	<u>WO 1997-EP5683</u>	19971015
W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG <u>AU 9748674</u> A1 19980515 <u>AU 1997-48674</u> 19971015 <u>PRIORITY APPLN. INFO.:</u> <u>DE 1996-19643331</u> A 19961021 <u>WO 1997-EP5683</u> W 19971015 OTHER SOURCE(S): MARPAT 128:294701 GI				



AB RaZNRbABD [I; A = Z1Z2; B = CO, CH2CO, OCH2CO, NHCH2CO, etc.; D = OH,
 (phenyl)alkoxy, cycloalkyloxy, etc.; Ra = H, (ar)alkyl, metabolically
 labile group, etc.; Rb = H, (cyclo)alkyl, aryl(alkyl), pyridyl(alkyl),
 ZRa, etc.; Z = 4,1'-bipiperidine-1,4'-diyl; Z1 = CO, CH2, CONH; Z2 =
 cyclohexylene, phenylene, etc.] were prepd. Thus, 4-(MeO)C6H4CH2NH2 was
 reductively condensed with 1-tert-butoxycarbonyl-4-piperidone and the
 product amidated by 4-(HO2C)C6H4OCH2CO2Me to give, in 3 addnl. steps,
 title compd. II. Data for biol. activity of I were given.

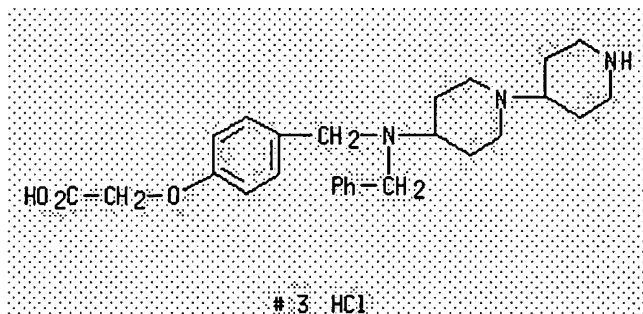
IT 206273-49-6P 206273-82-7P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of N-bipiperidinybenzamides and analogs as cell adhesion inhibitors)

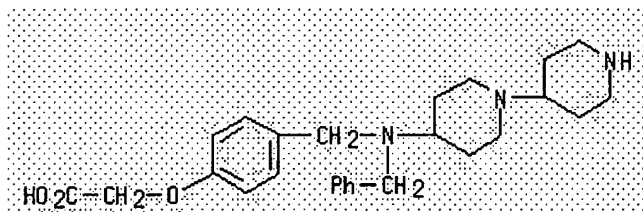
RN 206273-49-6 HCAPLUS

CN Acetic acid, [4-[[[1,4'-bipiperidin]-4-yl(phenylmethyl)amino]methyl]phenoxy]-, trihydrochloride (9CI) (CA INDEX NAME)



RN 206273-82-7 HCAPLUS

CN Acetic acid, [4-[[[1,4'-bipiperidin]-4-yl(phenylmethyl)amino]methyl]phenoxy]- (9CI) (CA INDEX NAME)



=> index PATENTS;s (JP 2000-369891)/pn;d rank

FILE 'ENCOMPPAT2' ACCESS NOT AUTHORIZED

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
28.03	221.40

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE	TOTAL
ENTRY	SESSION
-3.00	-3.75

CA SUBSCRIBER PRICE

INDEX 'CAOLD, CAPLUS, CASREACT, CROPU, DGENE, DPCI, ENCOMPPAT, EPFULL, FRANCEPAT, FRFULL, FSTA, GBFULL, IFIPAT, IMSPATENTS, INPADOC, JAPIO, KOREAPAT, LITALERT, NTIS, PAPERCHEM2, PATDD, PATDPA, PATDPAFULL, PATDPASPC, PCTFULL, PCTGEN, PIRA, PROUSDDR, PS, ...'

ENTERED AT 11:35:46 ON 02 FEB 2006

40 FILES IN THE FILE LIST IN STNINDEX

Enter SET DETAIL ON to see search term postings or to view search error messages that display as 0* with SET DETAIL OFF.

25 FILES SEARCHED...index PATENTS;s (JP 2000-369891)/pn;d rank

FILE 'ENCOMPPAT2' ACCESS NOT AUTHORIZED

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION

FULL ESTIMATED COST	28.03	221.40
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	-3.00	-3.75

INDEX 'CAOLD, CAPLUS, CASREACT, CROPU, DGENE, DPCI, ENCOMPPAT, EPFULL,
FRANCEPAT, FRFULL, FSTA, GBFULL, IFIPAT, IMSPATENTS, INPADOC, JAPIO,
KOREAPAT, LITALERT, NTIS, PAPERCHEM2, PATDD, PATDPA, PATDPAFULL,
PATDPASPC, PCTFULL, PCTGEN, PIRA, PROUSDDR, PS, ...'
ENTERED AT 11:35:46 ON 02 FEB 2006

40 FILES IN THE FILE LIST IN STNINDEX

Enter SET DETAIL ON to see search term postings or to view
search error messages that display as 0* with SET DETAIL OFF.

25 FILES SEARCHED...

0 FILES HAVE ONE OR MORE ANSWERS, 40 FILES SEARCHED IN STNINDEX

L13 QUE (JP 2000-369891)/PN

NO F-NUMBERS HAD GREATER THAN ZERO HITS

=> fil EPFULL;s (JP 2000-369891)/pn,apps

COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	1.22	222.62
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	0.00	-3.75

FILE 'EPFULL' ENTERED AT 11:36:54 ON 02 FEB 2006
COPYRIGHT (C) 2006 European Patent Office / FIZ Karlsruhe

FILE LAST UPDATED: 28 DEC 2005 <20051228/UP>
FILE COVERS 1978 TO DATE

>>> SIMULTANEOUS LEFT AND RIGHT TRUNCATION IS AVAILABLE
IN FIELDS /BI and /CLM. <<<

>>> UPDATING DELAYED DUE TO DELIVERY FORMAT CHANGES. <<<

>>> NEW IPC8 DATA AND FUNCTIONALITY NOT YET AVAILABLE IN THIS FILE.
USE IPC7 FORMAT FOR SEARCHING THE IPC. WATCH THIS SPACE FOR FURTHER
DEVELOPMENTS AND SEE OUR NEWS SECTION FOR FURTHER INFORMATION
ABOUT THE IPC REFORM <<<

0 (JP 2000-369891)/PN
(JP2000369891/PN)
0 JP2000-369891/AP
1 JP2000-369891/PRN
1 (JP 2000-369891)/APPS
(JP2000-369891/AP, PRN)

L14 1 (JP 2000-369891)/PN,APPS

=> d 1 imax

L14 ANSWER 1 OF 1 EPFULL COPYRIGHT 2006 EPO/FIZ KA on STN

Full
Text

ACCESSION NUMBER: 2001:108453 EPFULL
 DATA UPDATE DATE: 20040128
 DATA UPDATE WEEK: 200405
 TITLE (ENGLISH): PPAR (PEROXISOME PROLIFERATOR ACTIVATED RECEPTOR)
 ACTIVATORS
 TITLE (FRENCH): ACTIVEUR DE RECEPTEUR ACTIVE PAR LE PROLIFERATEUR DE
 PEROXISOME
 INVENTOR(S): YOSHIDA, Shinichi, 6-4-1, Nishifuna, Funabashi-shi,
 Chiba 273-0031, JP; SAKUMA, Shogo, 1-28-1-205,
 Yoshikawa, Yoshikawa-shi, Saitama 342-0055, JP; ENDO,
 Tsuyoshi, 3-23-6-A102, Wakagi, Itabashi-ku, Tokyo
 174-0065, JP; TENDO, Atsushi, 299-1-704, Oaza Hibori,
 Kasukabe-shi, Saitama 344-0005, JP; TAKAHASHI,
 Toshihiro, 4-1-6-505, Hikonari, Misato-shi, Saitama
 341-0033, JP; KOBAYASHI, Kunio, 5-1-3-202, Waseda,
 Misato-shi, Saitama 341-0018, JP; MOCHIDUKI, Nobutaka,
 1-30-238, Abiko, Abiko-shi, Chiba 270-1166, JP;
 YAMAKAWA, Tomio, 2-23-2, Midoridai, Kashiwa-shi, Chiba
 277-0884, JP; KANDA, Takashi, 3-10-18-701, Hikonari,
 Misato-shi, Saitama 341-0003, JP; MASUI, Seiichiro,
 4-6-21, Nakaduma, Ageo-shi, Saitama 362-0072, JP
 PATENT APPLICANT(S): NIPPON CHEMIPHAR CO., LTD., (CHEMIPHAR CO., LTD.,
 NIPPON), 2-3, Iwamoto-cho 2-chome, Chiyoda-ku, Tokyo
 101-8678, JP
 PATENT APPL. NUMBER: 706262
 LANGUAGE OF FILING: Japanese
 LANGUAGE OF PUBL.: English
 LANGUAGE OF PROCEDURE: English
 LANGUAGE OF TITLE: English; French
 DOCUMENT TYPE: Patent
 PATENT INFO TYPE: WOAI International application published with search
 report

PATENT INFORMATION:

	NUMBER	KIND	DATE
	WO 2002046176	A1	20020613
DESIGNATED STATES:	AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT		
	SE TR		
EXTENSION STATES:	AL LT LV MK RO SI		
APPLICATION INFO.:	EP 2001-999242	A	20011204
	WO 2001-JP10577	A	20011204
PRIORITY INFO.:	JP 2000-369891	A	20001205
INT. PATENT CLASSIF.:			
MAIN:	C07D277-56		
SECONDARY:	C07D277-68; C07D277-42; C07D277-82; C07D233-90; C07D233-88; C07D235-24; C07D235-30; A61K031-426; A61K031-428; A61K031-4164; A61K031-4184; A61P043-00		

LEGAL STATUS

AN 2001:108453 EPFULL
 20020807 WOB006EP The EPO has been informed by WIPO that EP was designated in
 this application

20020807 WOB870 PCT publication data
20020613
20020807 EPB840 Designated contracting states
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE TR
WO 2002046176 A1 20020613
20020807 EPB844EP Extension of the European patent to
AL LT LV MK RO SI
20020807 WOB880 PCT, Publication of the international search report (A3
publication)
20020613
20040128 WOB006EPR PCT application not entering the European phase
20040128 EPB237 Application deemed withdrawn
20030708

=> fil EPFULL;s (EP 2001-999242)/pn,apps

COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
5.41	228.03

FULL ESTIMATED COST

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE	TOTAL
ENTRY	SESSION
0.00	-3.75

CA SUBSCRIBER PRICE

FILE 'EPFULL' ENTERED AT 11:37:59 ON 02 FEB 2006

COPYRIGHT (C) 2006 European Patent Office / FIZ Karlsruhe

FILE LAST UPDATED: 28 DEC 2005

<20051228/UP>

FILE COVERS 1978 TO DATE

>>> SIMULTANEOUS LEFT AND RIGHT TRUNCATION IS AVAILABLE
IN FIELDS /BI and /CLM. <<<

>>> UPDATING DELAYED DUE TO DELIVERY FORMAT CHANGES. <<<

>>> NEW IPC8 DATA AND FUNCTIONALITY NOT YET AVAILABLE IN THIS FILE.
USE IPC7 FORMAT FOR SEARCHING THE IPC. WATCH THIS SPACE FOR FURTHER
DEVELOPMENTS AND SEE OUR NEWS SECTION FOR FURTHER INFORMATION
ABOUT THE IPC REFORM <<<

0 (EP 2001-999242)/PN
(EP2001999242/PN)

1 EP2001-999242/AP

0 EP2001-999242/PRN

1 (EP 2001-999242)/APPS

(EP2001-999242/AP, PRN)

L15

1 (EP 2001-999242)/PN,APPS

=> d 1 imax

L15 ANSWER 1 OF 1 EPFULL COPYRIGHT 2006 EPO/FIZ KA on STN

Full
Text

ACCESSION NUMBER: 2001:108453 EPFULL

DATA UPDATE DATE: 20040128

DATA UPDATE WEEK: 200405

TITLE (ENGLISH): PPAR (PEROXISOME PROLIFERATOR ACTIVATED RECEPTOR)
ACTIVATORS

TITLE (FRENCH): ACTIVATEURS DE RECEPTEUR ACTIVE PAR LE PROLIFERATEUR DE PEROXISOME

INVENTOR(S): YOSHIDA, Shinichi, 6-4-1, Nishifuna, Funabashi-shi, Chiba 273-0031, JP; SAKUMA, Shogo, 1-28-1-205, Yoshikawa, Yoshikawa-shi, Saitama 342-0055, JP; ENDO, Tsuyoshi, 3-23-6-A102, Wakagi, Itabashi-ku, Tokyo 174-0065, JP; TENDO, Atsushi, 299-1-704, Oaza Hibori, Kasukabe-shi, Saitama 344-0005, JP; TAKAHASHI, Toshihiro, 4-1-6-505, Hikonari, Misato-shi, Saitama 341-0033, JP; KOBAYASHI, Kunio, 5-1-3-202, Waseda, Misato-shi, Saitama 341-0018, JP; MOCHIDUKI, Nobutaka, 1-30-238, Abiko, Abiko-shi, Chiba 270-1166, JP; YAMAKAWA, Tomio, 2-23-2, Midoridai, Kashiwa-shi, Chiba 277-0884, JP; KANDA, Takashi, 3-10-18-701, Hikonari, Misato-shi, Saitama 341-0003, JP; MASUI, Seiichiro, 4-6-21, Nakaduma, Ageo-shi, Saitama 362-0072, JP

PATENT APPLICANT(S): NIPPON CHEMIPHAR CO., LTD., (CHEMIPHAR CO., LTD., NIPPON), 2-3, Iwamoto-cho 2-chome, Chiyoda-ku, Tokyo 101-8678, JP

PATENT APPL. NUMBER: 706262

LANGUAGE OF FILING: Japanese

LANGUAGE OF PUBL.: English

LANGUAGE OF PROCEDURE: English

LANGUAGE OF TITLE: English; French

DOCUMENT TYPE: Patent

PATENT INFO TYPE: WOAI International application published with search report

PATENT INFORMATION:

	NUMBER	KIND	DATE
	<u>WO 2002046176</u>	A1	20020613
DESIGNATED STATES:	AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE TR		
EXTENSION STATES:	AL LT LV MK RO SI		
<u>APPLICATION INFO.:</u>	<u>EP 2001-999242</u>	A	20011204
	<u>WO 2001-JP10577</u>	A	20011204
PRIORITY INFO.:	JP 2000-369891	A	20001205
INT. PATENT CLASSIF.:			
MAIN:	C07D277-56		
SECONDARY:	C07D277-68; C07D277-42; C07D277-82; C07D233-90; C07D233-88; C07D235-24; C07D235-30; A61K031-426; A61K031-428; A61K031-4164; A61K031-4184; A61P043-00		

LEGAL STATUS

AN 2001:108453 EPFULL

20020807 WOB006EP The EPO has been informed by WIPO that EP was designated in this application

20020807 WOB870 PCT publication data
20020613

20020807 EPB840 Designated contracting states
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE TR
WO 2002046176 A1 20020613

20020807 EPB844EP Extension of the European patent to
AL LT LV MK RO SI

20020807 WOB880 PCT, Publication of the international search report (A3 publication)
20020613

20040128 WOB006EPR PCT application not entering the European phase

20040128 EPB237 Application deemed withdrawn
20030708

=>